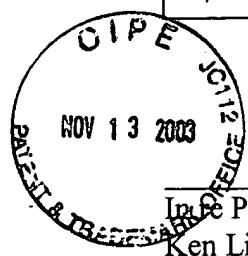


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Docket No.: 05432/000I004-US0
(PATENT)



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Patent Application of:
Ken Liljegren, et al.

Application No.: 09/730,380

Customer No.: 07278

Filed: December 5, 2000

Art Unit: 1625

For: PHARMACEUTICAL COMPOSITION
CONTAINING CITALOPRAM

Examiner: Charanjit AULAKH

INFORMATION DISCLOSURE STATEMENT (IDS)

MAIL BOX RCE
Commissioner for Patents
P.O. Box 1450
Alexandria, VA 22313-1450

Dear Sir:

Pursuant to 37 CFR 1.56, 1.97 and 1.98, attached hereto is a copy of Form PTO/SB/08 and copies of the documents listed thereon.

In accordance with MPEP Sections 609 and 707.05(b), it is requested that each document cited (including any cited in applicant's specification which is not repeated on the attached Form PTO/SB/08) be given thorough consideration and that it be cited of record in the prosecution history of the present application by initialing Form PTO/SB/08 next to the document. Such initialing is requested even if the Examiner does not consider a cited document to be sufficiently pertinent to use in a rejection, or otherwise does not consider it to be prior art for any reason, or even if the Examiner does not believe that the guidelines for citation have been fully complied with. This is requested so that each document becomes listed on the face of the patent issuing on the present application. This Information Disclosure Statement is filed before the mailing date of a first Office Action after the filing of a Request for Continued Examination under 37 CFR 1.114 (37 CFR 1.97(b)(4)).

A copy of each document on the PTO/SB/08 is attached.

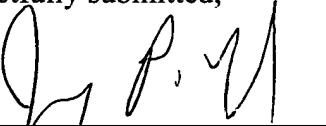
In accordance with 37 CFR 1.97(g), the filing of this Information Disclosure Statement shall not be construed to mean that a search has been made or that no other material information as defined in 37 CFR 1.56(a) exists. In accordance with 37 CFR 1.97(h), the filing of this Information Disclosure statement shall not be construed to be an admission that any patent, publication or other information referred to therein is "prior art" for this invention unless specifically designated as such.

It is submitted that the Information Disclosure Statement is in compliance with 37 CFR 1.98 and the Examiner is respectfully requested to consider the listed references.

The Commissioner is authorized to charge any deficiency of up to \$300.00 or credit any excess in this fee to Deposit Account No. 04-0100.

Dated: November 13, 2003

Respectfully submitted,

By 

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PTO/SB/08a/b (08-03)

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Substitute for form 1449A/B/PTO				Complete if Known	
				Application Number	09/730,380
				Filing Date	December 5, 2000
				First Named Inventor	Ken Liljegren
				Art Unit	1625
				Examiner Name	C. Aulakh
Sheet	1	of	2	Attorney Docket Number	05432/000I004-US0

U.S. PATENT DOCUMENTS					
Examiner Initials*	Cite No. ¹	Document Number	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		Number-Kind Code ² (if known)			
1	US-4,721,723	01-26-1988	Barnes et al.		
2	US-5,683,720	11-04-1997	Myers et al.		
3	US-5,840,334	11-24-1998	Raiden et al.		
4	US-5,869,098	02-09-1999	Misra et al.		
5	US-5,980,941	11-09-1999	Raiden et al.		

FOREIGN PATENT DOCUMENTS					
Examiner Initials*	Cite No. ¹	Foreign Patent Document	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		Country Code ³ -Number ⁴ -Kind Code ⁵ (if known)			
6	EP-07140663 A3		01-15-1997	Eli Lilly and Company	
7	GB-2357762		07-04-2001	H Lundbeck A/S	
8	GB-1358915		07-03-1974	Merck & Co, Inc.	
9	WO-99/03469		01-28-1999	Smithkline Beecham	
10	WO-01/68627		09-20-2001	H. Lundbeck A/S	
11	CA-2291072		05-14-1998	H. Lundbeck A/S	
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13	CA-2291067		05-14-1998	H. Lundbeck A/S	
14	CA-2178637		06-22-1995	Smithkline Beecham	
15	CA-2291134		04-20-2000	H. Lundbeck A/S	
16	CA-2163840		05-29-1996	Eli Lilly and Company	

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NON PATENT LITERATURE DOCUMENTS					
Examiner Initials	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.			T ²
	17	Webpage from Lundbeck website (www.lundbeck.com), company's activites			
	18	Webpage from Lundbeck website (www.lundbeck.com): Product information on Cipramil			
	19	Remington's Pharmaceutical Sciences, 18th Edition, Chapter 89, Oral Solid Dosage Forms, pp. 1633-1658			
	20	Bhogi B. Sheth, et al., Compressed Tablets, Chapter 3 in Pharmaceutical Dosage Forms: Tablets, Vol. 1, H. Lieberman and L. Lachman eds., Marcel Dekker, Inc., New York and Basel, 1979, pp. 109-185			
	21	Chapters 2 to 4 in Pharmaceutical Dosage Forms: Tablets, Vol. 1, H. Lieberman and L. Lachman, eds., Marcel Dekker, Inc. New York and Basel 1989, pp.75-246 (Chapter 2: Tablet			

{W:054 320001}	Date Considered
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PTO/SB/08a/b (08-03)

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Substitute for form 1449A/B/PTO				Complete if Known	
INFORMATION DISCLOSURE STATEMENT BY APPLICANT				Application Number	09/730,380
<i>(Use as many sheets as necessary)</i>				Filing Date	December 5, 2000
				First Named Inventor	Ken Liljegren
				Art Unit	1625
				Examiner Name	C. Aulakh
Sheet	2	of	2	Attorney Docket Number	05432/000I004-US0

		and Formulation Design; Chapter 3: Compressed Tablets by Wet Granulation; Chapter 4: Compressed Tablets by Direct Compression)	
	22	Keith Marshall, "Compression and Consolidation of Powdered Solids, Chapter 4, The Theory and Practise of Industrial Pharmacy, Lieberman, Lachman, and Kanig, eds., 3rd Edition, 1986, pp. 66-99	
	23	Hoener et al., Chapter 4, Factors Influencing Drug Absorption and Drug Availability, Modern Pharmaceutics, 3rd edition, Banker and Rhodes, eds., Marcel Dekker, New York and Basel, 1995, pp. 121-153	
	24	Edward M. Rudnic, et al., Chapter 10, Tablet Dosage Forms, Modern Pharmaceutics, 3d edition, Banker and Rhodes, eds., Marcel dekker, New York and Basel, 1995, pp. 333-394	
	25	Joseph B. Schwartz, et al., Chapter 18, Optimazation Techniques in Pharmaceutical Formulation and Processing, Modern Pharmaceutics, 3rd Edition, Banker and Rhodes, eds., Marcel Dekker, New York and Basel, 1995, pp. 727-752	
	26	Gunsel, et al, Chapter 11, Tablets, The Theory and Practice of Industrial Pharmacy, Lieberman, Lachman, and Kanig, eds., 2nd Edition, 1976,pp. 321-358	
	27	Keith Marshall, Chapter 10, Solid Oral Dosage Forms, Modern Pharmaceutics, 1st Edition, Banker and Rhodes, eds., Marcel Dekker, New York and Basel, 1979, p. 359-427	
	28	Vogel's Textbook of Practical Organic Chemistry, Fourth Edition, pp. 100-263	
	29	Dr. Fritz Gstirner, Professor fur Pharmazeutische Technologie an der Universitat Bonn, 1973, Einfuhrung in Die Verfahrenstechnik Der Arzneiformung, pp. 201-203	
	30	O'Connor, R.E. et al., Chapter 91 Powders, Remington: The Science and Practise of Pharmacy, 19th Ed., A. Gennaro, editor, Mack Publishing Co., Easton, 1995, pages 1598-1613	
	31	Banker, G.S., et al., Chapter 11, Tablets, The Theory and Practice of Industrial Pharmacy, Lieberman, Lachman, and Kanig, eds, 3rd Edition, 1986, pgs. 293-345	
	32	Hoener et al., Chapter 4, Factors Influencing Drug Absorption and Drug Availability, Modern Pharmaceutics, 1st edition, Banker and Rhodes, eds., Marcel Dekker, New York and Basel, 1979, pp. 143-182	
	33	Joseph B. Schwartz, et al., Chapter 17, Optimazation Techniques in Pharmaceutical Formulation and Processing, Modern Pharmaceutics, 1st Edition, Banker and Rhodes, eds., Marcel Dekker, New York and Basel, 1979, pp. 711-734	

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{W:1054 320001}	Date Considered	
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